

About the Author

Mr. Rizzo is an epidemiologist with the Healthcare-Associated Infections Program at the California Department of Public Health. His work focuses on surveillance of antimicrobial-resistant healthcare-associated infections and evaluation of prevention programs.

References

- Centers for Disease Control and Prevention. Antibiotic/antimicrobial resistance. Biggest threats and data. 2018 [cited 2019 Feb 19]. https://www.cdc.gov/drugresistance/biggest_threats.html
- Huang SS, Avery TR, Song Y, Elkins KR, Nguyen CC, Nutter SK, et al. Quantifying interhospital patient sharing as a mechanism for infectious disease spread. *Infect Control Hosp Epidemiol*. 2010;31:1160–9. <http://dx.doi.org/10.1086/656747>
- Centers for Disease Control and Prevention. Patient safety atlas. 2018 [cited 2018 Jul 13]. <https://www.cdc.gov/hai/surveillance/ar-patient-safety-atlas.html>
- Weiner LM, Webb AK, Limbago B, Dudeck MA, Patel J, Kallen AJ, et al. Antimicrobial-resistant pathogens associated with healthcare-associated infections: summary of data reported to the National Healthcare Safety Network at the Centers for Disease Control and Prevention, 2011–2014. *Infect Control Hosp Epidemiol*. 2016;37:1288–301. <http://dx.doi.org/10.1017/ice.2016.174>
- Woodworth KR, Walters MS, Weiner LM, Edwards J, Brown AC, Huang JY, et al. Vital signs: containment of novel multidrug-resistant organisms and resistance mechanisms—United States, 2006–2017. *MMWR Morb Mortal Wkly Rep*. 2018;67:396–401. <http://dx.doi.org/10.15585/mmwr.mm6713e1>
- National Healthcare Safety Network, Centers for Disease Control and Prevention. Surveillance reporting for enrolled facilities, 2015 [cited 2018 Jul 10]. <https://www.cdc.gov/nhsn/enrolled-facilities/index.html>
- Magiorakos A-P, Srinivasan A, Carey RB, Carmeli Y, Falagas ME, Giske CG, et al. Multidrug-resistant, extensively drug-resistant and pandrug-resistant bacteria: an international expert proposal for interim standard definitions for acquired resistance. *Clin Microbiol Infect*. 2012;18:268–81. <http://dx.doi.org/10.1111/j.1469-0691.2011.03570.x>
- Kadri SS, Adjemian J, Lai YL, Spaulding AB, Ricotta E, Prevots DR, et al.; National Institutes of Health Antimicrobial Resistance Outcomes Research Initiative (NIH-ARORI). Difficult-to-treat resistance in gram-negative bacteremia at 173 US hospitals: retrospective cohort analysis of prevalence, predictors, and outcome of resistance to all first-line agents. *Clin Infect Dis*. 2018;67:1803–14.
- Soe MM, Edwards JR, Sievert DM, Ricks PM, Magill SS, Fridkin KN. Evaluating state-specific antibiotic resistance measures derived from central line-associated bloodstream infections, national healthcare safety network, 2011. *Infect Control Hosp Epidemiol*. 2015;36:54–64. <http://dx.doi.org/10.1017/ice.2014.11>
- Talan DA, Takhar SS, Krishnadasan A, Abrahamian FM, Mower WR, Moran GJ; EMERGENCY ID Net Study Group. Fluoroquinolone-resistant and extended-spectrum β -lactamase-producing *Escherichia coli* infections in patients with pyelonephritis, United States. *Emerg Infect Dis*. 2016;22. <http://dx.doi.org/10.3201/eid2209.160148>
- Frazer BW, Trivedi T, Montgomery M, Petrovic D-F, Yamaji R, Riley L. Emergency department urinary tract infections caused by extended-spectrum β -lactamase-producing *Enterobacteriaceae*: many patients have no identifiable risk factor and discordant empiric therapy is common. *Ann Emerg Med*. 2018;72:449–56. <http://dx.doi.org/10.1016/j.annemergmed.2018.05.006>

Address for correspondence: Kyle Rizzo, California Department of Public Health, Healthcare-Associated Infections Program, 850 Marina Bay Pkwy, Building E, 1st Floor, Richmond, CA 94804, USA; email: kyle.rizzo@cdph.ca.gov

etymologia

Carbapenem [kahr''bē-pen'əm]

Ronnie Henry

A class of broad-spectrum β -lactam antibiotics, structurally similar to penicillins, with the substitution of a carbon atom (*carba*-) for a sulfur atom. This substitution creates a double bond on the pentane ring, which becomes a pentene ring (*-penem*).

The first carbapenem, thienamycin (*theion* ["sulfur"] + *enamine* [an unsaturated compound that forms the backbone of the molecule] + *-mycin* [suffix for drugs produced by *Streptomyces* spp.]), was discovered in 1976 in culture

broths of the newly recognized species *Streptomyces cattleya*. Thienamycin rapidly decomposes in the presence of water, which limits its clinical utility.

The first carbapenem approved for use in the United States was imipenem, the stable N-formimidoyl derivative of thienamycin, in 1985. Resistance to imipenem, encoded on a mobile genetic element, was first identified in *Pseudomonas aeruginosa* in Japan in 1991, and carbapenemase-producing organisms have since spread globally.

Sources

- Kahan JS, Kahan FM, Goegelman R, Currie SA, Jackson M, Stapley EO, et al. Thienamycin, a new beta-lactam antibiotic. I. Discovery, taxonomy, isolation and physical properties. *J Antibiot (Tokyo)*. 1979;32:1–12. <http://dx.doi.org/10.7164/antibiotics.32.1>
- Kesado T, Hashizume T, Asahi Y. Antibacterial activities of a new stabilized thienamycin, N-formimidoyl thienamycin, in comparison with other antibiotics. *Antimicrob Agents Chemother*. 1980;17:912–7. <http://dx.doi.org/10.1128/AAC.17.6.912>
- Meletis G. Carbapenem resistance: overview of the problem and future perspectives. *Ther Adv Infect Dis*. 2016;3:15–21. <http://dx.doi.org/10.1177/2049936115621709>

Address for correspondence: Ronnie Henry, Centers for Disease Control and Prevention, 1600 Clifton Rd NE, Mailstop E28, Atlanta, GA 30329-4027, USA; email: boq3@cdc.gov

DOI: <https://doi.org/10.3201/eid2507.ET2507>